

## Synthesis of new $\text{Ln}_4(\text{Al}_2\text{O}_6\text{F}_2)\text{O}_2$ ( $\text{Ln} = \text{Sm}, \text{Eu}, \text{Gd}$ ) phases with a cuspidine-related structure

Morán-Ruiz, Aroa; Wain-Martin, Aritza; Orera, Alodia; Sanjuán, Maria Luisa; Larrañaga, Aitor; Slater, Peter; Arriortua, Maribel

DOI:

[10.1107/S205225251801744X](https://doi.org/10.1107/S205225251801744X)

### Document Version

Publisher's PDF, also known as Version of record

### Citation for published version (Harvard):

Morán-Ruiz, A, Wain-Martin, A, Orera, A, Sanjuán, ML, Larrañaga, A, Slater, P & Arriortua, M 2019, 'Synthesis of new  $\text{Ln}_4(\text{Al}_2\text{O}_6\text{F}_2)\text{O}_2$  ( $\text{Ln} = \text{Sm}, \text{Eu}, \text{Gd}$ ) phases with a cuspidine-related structure', *IUCrJ*, vol. 6, no. Pt 1, pp. 128-135. <https://doi.org/10.1107/S205225251801744X>

[Link to publication on Research at Birmingham portal](#)

### General rights

Unless a licence is specified above, all rights (including copyright and moral rights) in this document are retained by the authors and/or the copyright holders. The express permission of the copyright holder must be obtained for any use of this material other than for purposes permitted by law.

- Users may freely distribute the URL that is used to identify this publication.
- Users may download and/or print one copy of the publication from the University of Birmingham research portal for the purpose of private study or non-commercial research.
- User may use extracts from the document in line with the concept of 'fair dealing' under the Copyright, Designs and Patents Act 1988 (?)
- Users may not further distribute the material nor use it for the purposes of commercial gain.

Where a licence is displayed above, please note the terms and conditions of the licence govern your use of this document.

When citing, please reference the published version.

### Take down policy

While the University of Birmingham exercises care and attention in making items available there are rare occasions when an item has been uploaded in error or has been deemed to be commercially or otherwise sensitive.

If you believe that this is the case for this document, please contact [UBIRA@lists.bham.ac.uk](mailto:UBIRA@lists.bham.ac.uk) providing details and we will remove access to the work immediately and investigate.

Received 28 August 2018

Accepted 10 December 2018

Edited by C.-Y. Su, Sun Yat-Sen University, China

**Keywords:** cuspidine-type rare-earth aluminates; fluorination; X-ray diffraction; poly(vinylidene difluoride); Raman spectroscopy.

**CCDC references:** 1886577; 1886578

**Supporting information:** this article has supporting information at [www.iucrj.org](http://www.iucrj.org)

# Synthesis of new $\text{Ln}_4(\text{Al}_2\text{O}_6\text{F}_2)\text{O}_2$ (Ln = Sm, Eu, Gd) phases with a cuspidine-related structure

Aroa Morán-Ruiz,<sup>a,\*</sup> Aritza Wain-Martin,<sup>a</sup> Alodia Orera,<sup>b</sup> María Luisa Sanjuán,<sup>b</sup> Aitor Larrañaga,<sup>a</sup> Peter R. Slater<sup>c</sup> and Maribel Arriortua<sup>a,d,\*</sup>

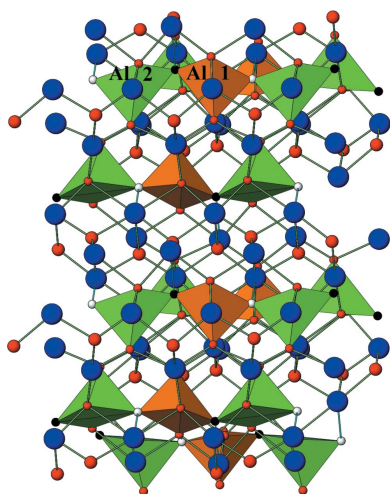
<sup>a</sup>Universidad del País Vasco (UPV/EHU), Facultad de Ciencia y Tecnología, Barrio Sarriena S/N, Leioa, Vizcaya 48940, Spain, <sup>b</sup>Instituto de Ciencia de Materiales de Aragón (CSIC - Universidad de Zaragoza), C/ Pedro Cerbuna 12, Zaragoza 50009, Spain, <sup>c</sup>University of Birmingham, School of Chemistry, Birmingham B15 2TT, UK, and <sup>d</sup>BCMaterials (Basque Centre for Materials, Applications and Nanostructures), Bld. Martina Casiano, 3rd. Floor, UPV/EHU Science Park, Barrio Sarriena S/N, Leioa, Vizcaya 48940, Spain. \*Correspondence e-mail: [aroa.moran@ehu.eus](mailto:aroa.moran@ehu.eus), [maribel.arriortua@ehu.eus](mailto:maribel.arriortua@ehu.eus)

The first fluorination of the cuspidine-related phases of  $\text{Ln}_4(\text{Al}_2\text{O}_7\Box)\text{O}_2$  (where Ln = Sm, Eu, Gd) is reported. A low-temperature reaction with poly(vinylidene difluoride) lead to the fluorine being substituted in place of oxygen and inserted into the vacant position between the dialuminate groups. X-ray photoelectron spectroscopy shows the presence of the F 1s photoelectron together with an increase in Al 2p and rare-earth 4d binding energies supporting F incorporation. Energy-dispersive X-ray spectroscopy analyses are consistent with the formula  $\text{Ln}_4(\text{Al}_2\text{O}_6\text{F}_2)\text{O}_2$ , confirming that substitution of one oxygen by two fluoride atoms has been achieved. Rietveld refinements show an expansion in the cell upon fluorination and confirm that the incorporation of fluoride in the  $\text{Ln}_4(\text{Al}_2\text{O}_7\Box)\text{O}_2$  structure results in changes in Al coordination from four to five. Thus, the isolated tetrahedral dialuminate  $\text{Al}_2\text{O}_7$  groups are converted to chains of distorted square-based pyramids. These structural results are also discussed based on Raman spectra.

## 1. Introduction

Minerals belonging to the cuspidine group have the general stoichiometry  $M_4(\text{Si}_2\text{O}_7)X_2$  ( $M$  = divalent cation;  $X$  = OH, F, O), with  $\text{Ca}_4(\text{Si}_2\text{O}_7)(\text{OH},\text{F})_2$  being the archetype compound. The cuspidine structure can be described as built up of chains of edge-sharing  $\text{MO}_7/\text{MO}_8$  polyhedra running parallel to the  $a$  axis (in the  $P2_1/c$  space group); the tetrahedral disilicate groups ( $\text{Si}_2\text{O}_7$ ) interconnect with these ribbons through the vertices. The structural formula of cuspidine is better described as  $\text{Ca}_4(\text{Si}_2\text{O}_7\Box)(\text{OH},\text{F})_2$  to directly show the vacant position between the disilicate groups. The filling of that position may convert the isolated pyrogro groups into infinite chains of distorted trigonal bipyramids (Martín-Sedeño *et al.*, 2004).

Other systems also adopt this structural type, including the  $\text{Ln}_4(\text{Al}/\text{Ga})_2\text{O}_9$  (Ln = rare-earth) type phases, which have attracted attention because of their ionic conductivity and thermal stability (Ghosh, 2015; Zhou *et al.*, 2014; Martín-Sedeño *et al.*, 2006; Morán-Ruiz *et al.*, 2018). In more recent years, the preparation and characterization of inorganic oxyfluorides have attracted significant interest. Thus, low-temperature fluorination methods can alter chemistry of the precursor oxide in different ways by charge compensation effects (Clemens & Slater, 2013). In particular, polymer reagents such as poly(vinylidene fluoride) and poly(tetrafluoroethylene) have been proven to be successful low-temperature fluorinating reagents, following the early work by



OPEN ACCESS

Slater (2002) which illustrates the use of PVDF to prepare  $\text{Ca}_2\text{CuO}_2\text{F}_2$  and  $\text{Sr}_2\text{TiO}_3\text{F}_2$ . Since then, a wide range of perovskite and related phases have been successfully fluorinated using this polymer route (Clemens *et al.*, 2014; Hancock *et al.*, 2012; Berry *et al.*, 2008; Heap *et al.*, 2007), and the method has been shown to be equally applicable to the fluorination of thin films (Kawahara *et al.*, 2017; Katayama *et al.*, 2016; Moon *et al.*, 2015). This earlier research has mainly focused on the fluorination of transition-metal containing materials, and so here we investigate the potential use for the fluorination of oxide systems that do not contain transition metals. In particular, given the recent studies on oxide ion/proton conductivity in  $\text{La}_4(\text{Ga}_{2-x}\text{Ti}_x\text{O}_{7+x/2})\text{O}_2$ , which illustrate the ability of the cuspidine structure to accommodate extra anions (Martín-Sedeño *et al.*, 2005), this would appear to be an ideal structure to examine the possible incorporation of fluoride. We have therefore investigated the fluorination of  $\text{Ln}_4(\text{Al}_2\text{O}_7\Box)\text{O}_2$  to give new  $\text{Ln}_4\text{Al}_2\text{O}_{9-x}\text{F}_{2x}$  ( $\text{Ln} = \text{Sm}, \text{Eu}, \text{Gd}$ ) ( $0 \leq x \leq 1$ ) phases. Here we report the results of these first low-temperature fluorination reactions of a range of rare-earth aluminate cuspidine-related phases. The introduction of fluorine ( $2\text{F}^-$  replacing  $\text{O}^{2-}$ ) was achieved through a reaction with poly(vinylidene fluoride) (PVDF) as the fluorinating agent. We investigate the success and effects of fluorination on the starting structure by X-ray diffraction (XRD), X-ray photoelectron spectroscopy (XPS),  $^{27}\text{Al}$  solid-state nuclear magnetic resonance (NMR), Raman spectroscopy, scanning electron microscopy (SEM) and energy dispersive X-ray spectroscopy (EDX). The thermal stability of these samples after fluorination was evaluated in air through thermogravimetric analysis (TGA).

## 2. Experimental

### 2.1. Powder preparation

Starting precursor oxides of  $\text{Ln}_4(\text{Al}_2\text{O}_7\Box)\text{O}_2$  ( $\text{Ln} = \text{Sm}, \text{Eu}, \text{Gd}$ ) were prepared by the glycine nitrate combustion route using the appropriate quantities of metals and combustible substance as previously reported by Morán-Ruiz *et al.* (2018). The introduction of fluorine ( $2\text{F}^-$  replacing  $\text{O}^{2-}$ ) into the  $\text{Ln}_4(\text{Al}_2\text{O}_7\Box)\text{O}_2$  structure was achieved through a low-temperature ( $400^\circ\text{C}$ ) reaction with PVDF (Slater, 2002) as the fluorinating agent. Thus, fluorination was achieved by mixing the rare-earth aluminate phase with PVDF in a 1:1 mol ratio (precursor oxide:  $\text{CH}_2\text{CF}_2$  monomer unit) and heating ( $80^\circ\text{C h}^{-1}$ ) the mixture at  $400^\circ\text{C}$  for 12 h in air.

Since poly(tetrafluoroethylene) (PTFE) has also been shown to be a very good fluorinating reagent, we investigated the possibility of fluorination of  $\text{Eu}_4(\text{Al}_2\text{O}_7\Box)\text{O}_2$  with PTFE under the same conditions. This gave similar results to the reaction with PVDF, with an observed expansion in the unit cell consistent with F incorporation.

### 2.2. Characterization techniques

X-ray powder diffraction patterns were recorded with a Philips X'Pert-Pro diffractometer using graphite-mono-

chromated  $\text{Cu K}\alpha_{1,2}$  radiation ( $\lambda_1 = 1.5406 \text{ \AA}$ ;  $\lambda_2 = 1.5443 \text{ \AA}$ ). The compounds were scanned between  $15$  and  $90^\circ$  ( $2\theta$ ) in  $0.02^\circ$  steps, counting 380 s per step. In addition, a Bruker D8 Advance Vario diffractometer, equipped with a primary monochromator and a solid SolX detector, with energy discrimination optimized for such radiation ( $\text{Cu K}\alpha_1$ ,  $\lambda_1 = 1.5406 \text{ \AA}$ ), were also used to improve the quality of the XRD data for structure refinement. The overall measuring time was  $\sim 120$  h per pattern to have good statistics over the  $2\theta$  angular range of  $5$ – $100^\circ$  with a  $0.02^\circ$  step size. The fitting of the measured and calculated pattern structure refinement was carried out using the program *FullProf* (Rodríguez-Carvajal, 2011). Moreover, *Atoms62* software (Shape Software, 2005) was also used to illustrate the structure.

X-ray photoelectron spectroscopy (XPS) measurements were performed using an XPS spectrometer (SPECS). All XPS spectra were acquired using a monochromatic X-ray source producing  $\text{Al K}\alpha$  radiation ( $h\nu = 1486.6 \text{ eV}$ ) and recorded using a Specs Phoibos 150 analyser. An initial analysis of the elements present in the sample was carried out (wide scan: step energy  $1 \text{ eV}$ , dwell time  $0.1 \text{ s}$ , pass energy  $80 \text{ eV}$ ) and individual high-resolution spectra were obtained (detail scan: step energy  $0.1 \text{ eV}$ , dwell time  $0.1 \text{ s}$ , pass energy  $30 \text{ eV}$ ) with an electron take-off angle of  $90^\circ$ . The binding energies (BEs) were calibrated using the C  $1s$  peak ( $\text{BE} = 284.6 \text{ eV}$ ) as an internal standard. The spectra were fitted by *CasaXPS* 2.3.16 software, modelling the properly weighted sum of Gaussian and Lorentzian component curves, after background subtraction according to Shirley.

The  $^{27}\text{Al}$  solid-state NMR spectra were recorded on a Bruker Avance III, at  $9.4 \text{ T}$  under magic angle (MAS) at  $14 \text{ kHz}$  using a Bruker probe head  $4 \text{ mm MAS DVT X/Y/H}$ . The  $^{27}\text{Al}$  MAS NMR were recorded at  $104.27 \text{ MHz}$  using a single-pulse sequence with a  $4 \mu\text{s}$  rf pulse ( $\pi/2$ ); the relaxation delay was  $0.5 \text{ s}$  and a total of  $20\,000$  scans were accumulated. The  $^{27}\text{Al}$  chemical shifts were calibrated indirectly with  $\text{Al}(\text{NO}_3)_3$ .

For Raman scattering measurements a DILOR XY spectrometer with a CCD detector and  $2 \text{ cm}^{-1}$  of spectral resolution was used. The  $514.5 \text{ nm}$  line of an  $\text{Ar}^+$ -ion laser was used as the excitation source, and the power output was kept below  $20 \text{ mW}$  after verifying that no changes were induced in the samples. A  $50\times$  microscope objective lens was used both for excitation and dispersed light collection. Some spectra were also collected in a WITEC Alpha 300M+ spectrometer working with  $633 \text{ nm}$  excitation. For each material, at least 3–4 representative spectra of different sample zones were recorded.

Thermogravimetric analyses were performed for all compositions on a TA Instruments SDT 2960 simultaneous DSC–TGA balance. The temperature was varied from room temperature up to  $900^\circ\text{C}$  at a heating rate of  $3^\circ\text{C min}^{-1}$  in air.

Compositional analysis was performed using an analytical scanning electron microscope (SEM, JEOL JSM-7000 F) with an electron microanalysis probe EDX (Oxford Pentafet energy dispersive X-ray analyzer). Samples were coated with a coal graphite layer ( $10 \text{ nm}$ ) deposited by evaporation

(Quorum Q150T Sputter Coater) to provide electrical conductivity. Back-scattered electrons were measured at a 20 kV accelerating voltage and  $5 \times 10^{-9}$  A current. A measurement time of 100 s per point was established for data acquisition. EDX system calibration was performed by measuring the beam current on  $\text{Ln}_4(\text{Al}_2\text{O}_7\Box)\text{O}_2$  and  $\text{AlF}_3$  as standards, to allow quantitative elemental analyses. The data processing was performed using Oxford *Inca* software. The characteristic emission lines used for the analysis were  $K\alpha$  for Al and F, and  $M\alpha$  for Sm, Eu and Gd. The morphologies of the powders were observed using secondary electrons at an accelerating voltage of 20 kV, a current of  $1.1 \times 10^{-11}$  A and a working distance of 9 mm. These samples were metallized by gold sputtering for better image definition.

### 3. Results and discussion

The X-ray powder diffraction patterns recorded from  $\text{Ln}_4(\text{Al}_2\text{O}_7\Box)\text{O}_2$  (Ln = Sm, Eu, Gd) and their new fluorinated derivatives are shown in Fig. 1. The XRD patterns show that all the samples consist of a single phase without impurities. Moreover, the fluorination induces a shift in peak position to lower angles corresponding to an increase in unit-cell sizes as the total anion content increases.

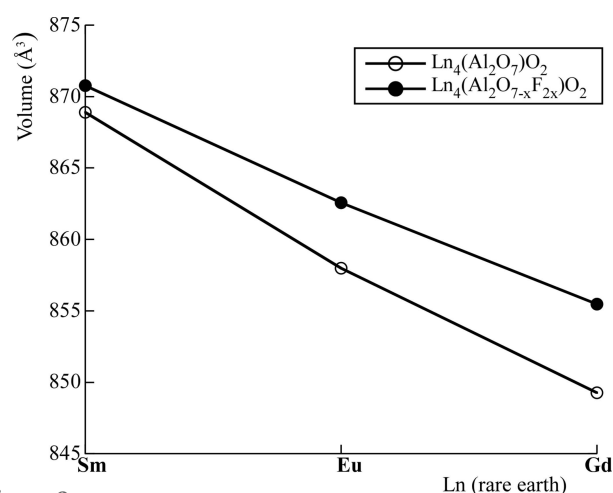
The volumes recorded from the pure oxides and their fluorinated derivatives are graphically represented in Fig. 2. From the data in the graphic it can be seen that the fluorination leads to a significant increase in unit-cell parameters. The volume difference between the starting oxide and fluorinated oxides becomes more noticeable as the rare earth size decreases. Moreover, the cell parameters change in good agreement with the variation of the ionic radii of the rare-earth cations, with the largest cell volume observed for the Sm system and the smallest for the Gd system (Morán-Ruiz *et al.*, 2018) [ $\text{Gd}^{3+}$  (coordination number VII): 1.00 Å;  $\text{Gd}^{3+}$  (VIII): 1.05 Å;  $\text{Eu}^{3+}$  (VII): 1.01 Å;  $\text{Eu}^{3+}$  (VIII): 1.07 Å;  $\text{Sm}^{3+}$  (VII): 1.02 Å;  $\text{Sm}^{3+}$  (VIII): 1.08 Å].

Representative SEM micrographs of the powder samples (as prepared and after fluorination at 400°C) are shown in Fig. 3. As observed, no significant differences can be seen in the morphology or the average particle size of the different

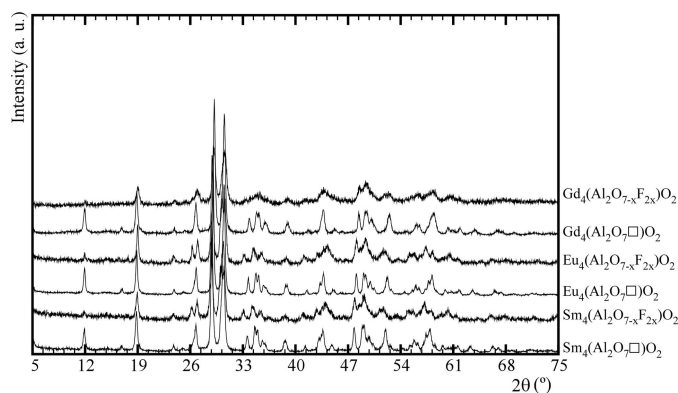
samples in these images. All samples are composed of agglomerated sub-micrometre particles.

The chemical compositions of the obtained fluorinated oxides were analysed using SEM-EDX. The measured values of the elements were checked on different points to obtain the average composition. The atomic percentage concentrations of detected elements are listed in Table 1. For comparison, data were also collected for  $\text{Eu}_4(\text{Al}_2\text{O}_7\Box)\text{O}_2$  fluorinated with half the molar equivalents of PVDF, in order to illustrate that F content can be controlled by the amount of polymer added.

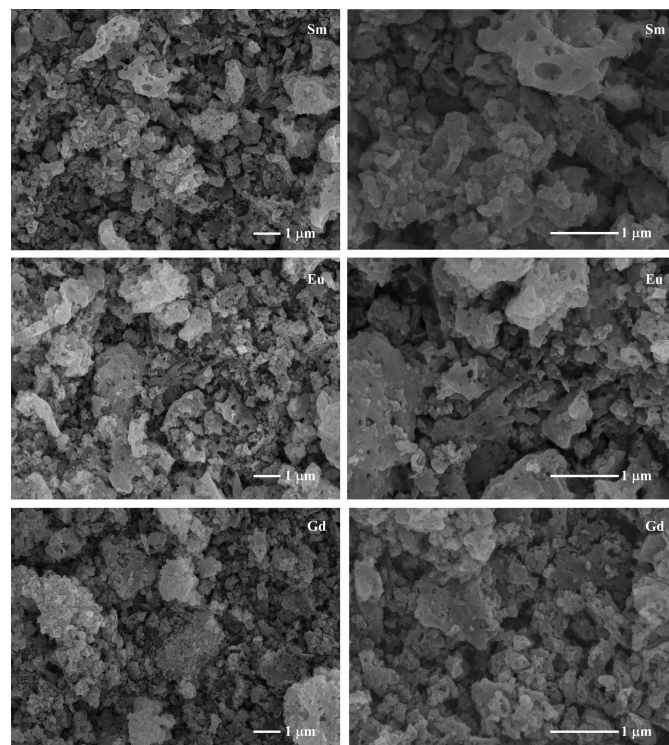
These results indicate that the substitution of two fluorine atoms for one oxygen is satisfactorily achieved to obtain new



**Figure 2**  
The volume changes between the pure oxides and their fluorinated derivatives.



**Figure 1**  
X-ray powder diffraction patterns recorded from materials of composition  $\text{Ln}_4(\text{Al}_2\text{O}_7\Box)\text{O}_2$  (Ln = Sm, Eu, Gd) and their fluorinated derivatives.



**Figure 3**  
Micrographs of  $\text{Sm}_4(\text{Al}_2\text{O}_6\text{F}_2)\text{O}_2$ ,  $\text{Eu}_4(\text{Al}_2\text{O}_6\text{F}_2)\text{O}_2$  and  $\text{Gd}_4(\text{Al}_2\text{O}_6\text{F}_2)\text{O}_2$  phases prepared using a low-temperature fluorination route.



**Table 1**

Chemical compositions (at.%) of fluorinated oxides obtained under various synthetic conditions.

The use of plasma-cleaning could reduce the fluorine content near to 12.5 at. %.

Sample	Ln <sup>†</sup>	Al	F
Sm <sub>4</sub> (Al <sub>2</sub> O <sub>7-x</sub> F <sub>2x</sub> )O <sub>2</sub>	24.0 (2)	12.2 (2)	13.4 (3)
Eu <sub>4</sub> (Al <sub>2</sub> O <sub>7-x</sub> F <sub>2x</sub> )O <sub>2</sub>	25.0 (4)	12.2 (3)	13.1 (1)
Gd <sub>4</sub> (Al <sub>2</sub> O <sub>7-x</sub> F <sub>2x</sub> )O <sub>2</sub>	25.6 (2)	12.6 (3)	13.6 (2)
Eu <sub>4</sub> (Al <sub>2</sub> O <sub>7-x</sub> F <sub>2x</sub> )O <sub>2</sub> ‡	23.2 (1)	12.0 (2)	6.9 (1)
Eu <sub>4</sub> (Al <sub>2</sub> O <sub>7-x</sub> F <sub>2x</sub> )O <sub>2</sub> §	22.7 (2)	11.5 (1)	12.7 (2)
Eu <sub>4</sub> (Al <sub>2</sub> O <sub>7-x</sub> F <sub>2x</sub> )O <sub>2</sub> ¶	23.0 (1)	11.5 (1)	12.3 (1)
Theoretical Ln <sub>4</sub> (Al <sub>2</sub> O <sub>6</sub> F <sub>2</sub> )O <sub>2</sub>	25	12.5	12.5

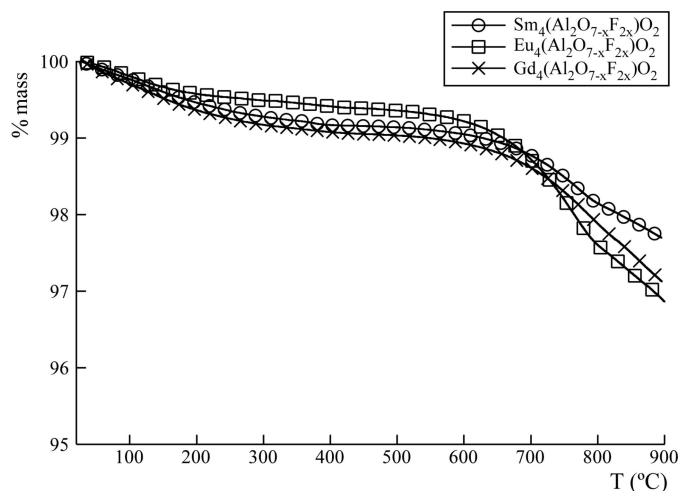
† Ln = Sm, Eu, Gd. ‡ Fluorination reaction using  $\frac{1}{2}$  PVDF [equivalent to 1 F ( $x = 0.5$ ) incorporation]. § Fluorination reaction using poly(tetrafluoroethylene) (PTFE). ¶ Fluorination reaction using  $\frac{1}{2}$  PTFE.

Ln<sub>4</sub>Al<sub>2</sub>O<sub>8</sub>F<sub>2</sub> (Ln = Sm, Eu, Gd) compositions. Examination of the fluorination with higher levels of PVDF led to no further increase in cell volume, illustrating the maximum F content had been reached. From these results it can be concluded that these cuspidine phases permit a maximum of two fluorine atoms per formula.

The samples were heated in a thermogravimetric analyzer in air at 900°C. The thermograms of all oxyfluorides are shown in Fig. 4. A decrease in mass with increasing temperature occurs between 550 and 900°C, which is associated with the loss of fluorine content due to the reaction with moisture in the air, leading to loss of HF and replacement by oxygen to reform the simple oxide system.

For all compositions, a gravimetric mass loss of ~3% is observed. From these results the (O/F)<sub>z</sub> relation is calculated (Table 2).

From the TGA data it can be concluded that the mass loss is not complete due to the low kinetic decomposition of these compounds. Preliminary studies show that the stabilization of the mass requires a long heating time (~6 h) at 1000°C (see supporting information, Fig. S1). In order to obtain the total



**Figure 4**  
Thermogravimetric analysis of new Ln<sub>4</sub>(Al<sub>2</sub>O<sub>6</sub>F<sub>2</sub>)O<sub>2</sub> (Ln = Sm, Eu, Gd) phases.

**Table 2**

Fluorine content loss calculated from the gravimetric mass loss (550–900°C).

Sample	x
Sm <sub>4</sub> (Al <sub>2</sub> O <sub>7-x</sub> F <sub>2x</sub> )O <sub>2</sub>	0.52
Eu <sub>4</sub> (Al <sub>2</sub> O <sub>7-x</sub> F <sub>2x</sub> )O <sub>2</sub>	0.93
Gd <sub>4</sub> (Al <sub>2</sub> O <sub>7-x</sub> F <sub>2x</sub> )O <sub>2</sub>	0.72

**Table 3**

Chemical compositions (at.%) of the TGA residues of fluorinated oxides.

Residue	Ln <sup>†</sup> (at.%)	Al (at.%)	F (at.%)
Sm <sub>4</sub> (Al <sub>2</sub> O <sub>7-x</sub> F <sub>2x</sub> )O <sub>2</sub>	24.8 (3)	12.6 (3)	6.6 (3)
Eu <sub>4</sub> (Al <sub>2</sub> O <sub>7-x</sub> F <sub>2x</sub> )O <sub>2</sub>	26.9 (1)	12.7 (1)	1.0 (1)
Gd <sub>4</sub> (Al <sub>2</sub> O <sub>7-x</sub> F <sub>2x</sub> )O <sub>2</sub>	27.4 (3)	11.5 (3)	2.8 (6)

† Ln = Sm, Eu, Gd.

**Table 4**

XPS analysis results of detected elements for the surface of the obtained Ln<sub>4</sub>(Al<sub>2</sub>O<sub>7-x</sub>F<sub>2x</sub>)O<sub>2</sub> compositions.

Sample	Ln 4d <sup>†</sup> (BE, eV)	Al 2p (BE, eV)	F 1s (BE, eV)
Sm <sub>4</sub> (Al <sub>2</sub> O <sub>7</sub> )O <sub>2</sub>	131.7	73.1	–
Sm <sub>4</sub> (Al <sub>2</sub> O <sub>7-x</sub> F <sub>2x</sub> )O <sub>2</sub>	133.1	74.3	685.0
Eu <sub>4</sub> (Al <sub>2</sub> O <sub>7</sub> )O <sub>2</sub>	135.5	73.1	–
Eu <sub>4</sub> (Al <sub>2</sub> O <sub>7-x</sub> F <sub>2x</sub> )O <sub>2</sub>	137.0	74.3	686.5
Gd <sub>4</sub> (Al <sub>2</sub> O <sub>7</sub> )O <sub>2</sub>	141.9	73.1	–
Gd <sub>4</sub> (Al <sub>2</sub> O <sub>7-x</sub> F <sub>2x</sub> )O <sub>2</sub>	142.8	74.3	685.3

† Ln = Sm, Eu, Gd.

fluorine content remaining in each sample after treatment at 900°C, the residues were analyzed by EDX. The atomic percentage concentrations of detected elements are summarized in Table 3. The obtained data coincide with the calculated fluorine content loss.

The success of the fluorination of rare-earth aluminates is also confirmed by XPS. A clear peak is observed in the analysed areas of the fluorinated oxides using a wide scan up to 1380 eV, attributable to an F 1s photoelectron (Fig. 5).

The resultant peak BEs before and after fluorination are presented in Table 4. In particular, we observe that fluorine incorporation induces an increase of the BEs of Al 2p and Ln 4d due to fluorine having a higher electronegativity than oxide.

This indicates greater electron transfer to fluorine, causing a decrease in the electron density at the cation and resulting in higher binding energy of the electrons from the core level of the cation (Dae-Min *et al.*, 2011). These peak-position shifts are observed in the high-resolution spectra of the Al 2p and Ln 4d spectral regions (Fig. 6).

The <sup>27</sup>Al NMR spectra of the fluorinated samples and Ln<sub>4</sub>(Al<sub>2</sub>O<sub>7</sub>)O<sub>2</sub> (Ln = Sm, Eu, Gd) are provided as supporting information (Fig. S2). It seems that the shape of the Ln<sub>4</sub>(Al<sub>2</sub>O<sub>6</sub>F<sub>2</sub>)O<sub>2</sub> spectra changes compared with the Ln<sub>4</sub>(Al<sub>2</sub>O<sub>7</sub>)O<sub>2</sub> spectra, which could be due to a modification of the coordination environment of Al<sup>3+</sup> in the fluorinated derivatives. However, the obtained <sup>27</sup>Al NMR data are not conclusive due to the paramagnetism of Sm, Eu and Gd rare-earth metals.

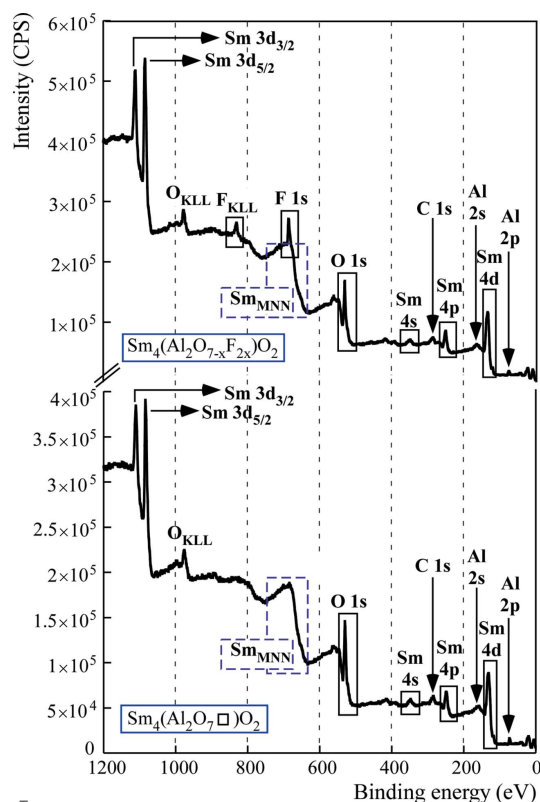


Figure 5

As an example, XPS survey spectra of the surface composition  $\text{Sm}_4(\text{Al}_2\text{O}_7\Box)\text{O}_2$  and their new fluorinated derivative.

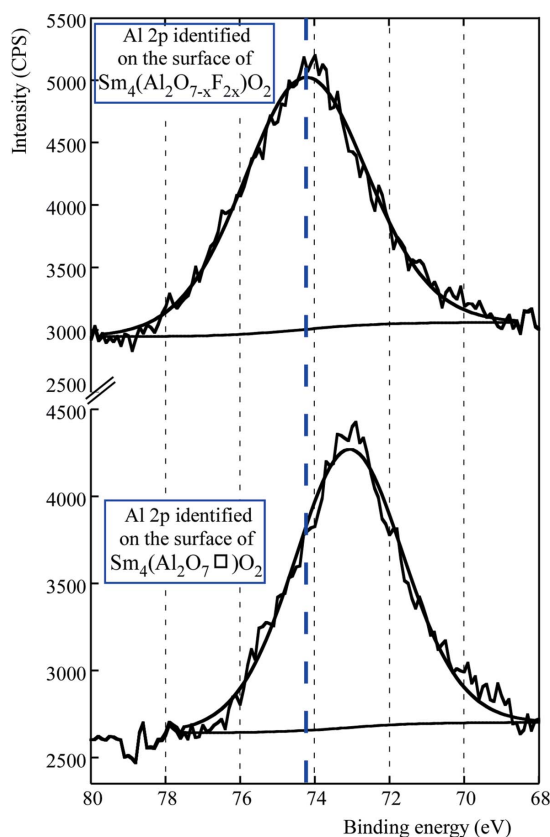


Figure 6

As an example, Al 2p spectral regions of the surface of  $\text{Sm}_4(\text{Al}_2\text{O}_7\Box)\text{O}_2$  and their new fluorinated derivative showing a shift to higher binding energy upon fluorination.

Full structural refinements of XRD data for  $\text{Sm}_4\text{Al}_2\text{O}_{9-x}\text{F}_{2x}$  and  $\text{Eu}_4\text{Al}_2\text{O}_{9-x}\text{F}_{2x}$  were carried out in the space group  $P2_1/c$  by using the  $\text{Sm}_4(\text{Al}_2\text{O}_7)\text{O}_2$  and  $\text{Eu}_4(\text{Al}_2\text{O}_7)\text{O}_2$  structures as starting models, respectively. Refined cell and positional parameters, obtained bond distances and angles, and the bond valences are summarized in Tables S1–S11. The Rietveld fittings of the X-ray data are displayed in Fig. 7.

After the convergence of the overall parameters, the occupation of the bridge oxygen site O(5) was replaced by F(1) (Kendrick *et al.*, 2008) and an extra fluorine position, F(2) (Martín-Sedeño *et al.*, 2006), was added in the vacant anion site between two  $\text{Al}_2\text{O}_7$  units, in order to account for the increase in anion content, and then refined. In both refinements, bond-length constraints were applied. The quality factors of the refinements are given in Table 5. It should be noted that distinguishing O and F by either X-ray or neutron diffraction is very difficult because of the nearly identical scattering factors. Therefore, the respective positions are commonly inferred by bond valence sum (BVS) calculations from the determined bond distances. In this respect, neutron diffraction data would lead to more accurate O/F positions and hence bond distances. However, Gd, Sm and Eu all show very strong neutron absorption, which makes such studies impractical. Therefore we have used BVS calculations based on the structures determined from the X-ray diffraction data.

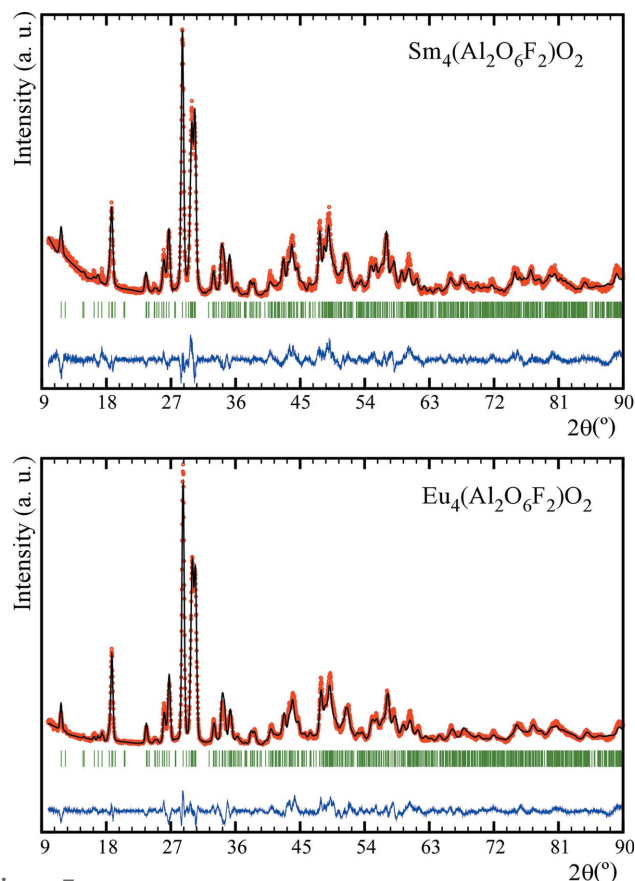


Figure 7

Rietveld refinement for new  $\text{Sm}_4(\text{Al}_2\text{O}_6\text{F}_2)\text{O}_2$  and  $\text{Eu}_4(\text{Al}_2\text{O}_6\text{F}_2)\text{O}_2$  cuspidine-related materials.

Table 5

The quality of refinements performed on new fluorinated oxides.

Samples	$\text{Sm}_4\text{Al}_2\text{O}_{9-x}\text{F}_{2x}$	$\text{Eu}_4\text{Al}_2\text{O}_{9-x}\text{F}_{2x}$
$\chi^2$	3.7	5.7
$R_{\text{Bragg}}$	12.8	11.7
$R_{\text{f}}$	10.6	9.2
$R_{\text{p}}$	10.9	10.2
$R_{\text{wp}}$	13.6	12.8
$R_{\text{exp}}$	7.06	5.39

These calculations are in agreement with the assignment of the F positions proposed, which is further supported by the Raman results (see later). In addition, the BVS values that are calculated for the F(1) and F(2) sites, assuming O is present, show a critical deficit of valence charge in the oxygen atoms. These results add further weight to the conclusion that F occupies these sites.

The introduction of fluorine leads to the conversion of isolated  $\text{M}_2\text{O}_7\Box$  groups into infinite chains of distorted square-based pyramids along the  $a$  axis, as observed for  $\text{La}_4(\text{Ti}_2\text{O}_8)\text{O}_2$ .

It is interesting to compare the present results with those of Si cuspidines of the  $\text{M}_4(\text{Si}_2\text{O}_7)\text{F}_2$  type (Achary *et al.*, 2017), where the  $\text{Si}_2\text{O}_7$  units are preserved and fluorine occupies the O(8) and O(9) sites instead of filling the anionic vacancies and substituting for bridge oxygen ions along the  $\text{AlO}_4$  chains. The different behaviour can be attributed to the larger size of Al cations compared with Si, and its higher ability to accommodate coordination numbers greater than four.

In summary, the structures of  $\text{Ln}_4(\text{Al}_2\text{O}_6\text{F}_2)\text{O}_2$  ( $\text{Ln} = \text{Sm}, \text{Eu}$ ) are monoclinic ( $P2_1/c$ ) with two sites for fluorine between the aluminate groups. Thus, as observed from Figs. 8 and 9, the aluminium coordination changes from four to five. Because of low crystallinity, the  $\text{Gd}_4(\text{Al}_2\text{O}_{7-x}\text{F}_x)\text{O}_2$  diffractogram produces a poor signal, which limits its Rietveld refinement. This lower crystallinity is probably related to the fact that Gd is the smallest rare-earth metal and also the large volume change upon fluorination, which may have reduced the particle size/crystallinity. Considering  $\text{Ln}_4(\text{Al}_2\text{O}_6\text{F}_2)\text{O}_2$  ( $\text{Ln} = \text{Sm}, \text{Eu}$ ) as representative structures of the obtained  $\text{Ln}_4(\text{Al}_2\text{O}_6\text{F}_2)\text{O}_2$  ( $\text{Ln} = \text{Sm}, \text{Gd}$ ) compositions, similar results could be expected for the gadolinium sample.

These structural assumptions have been further discussed based on Raman results. Raman spectra are shown in Fig. 10 for samples  $\text{Ln}_4(\text{Al}_2\text{O}_7\Box)\text{O}_2$  (bottom set) and  $\text{Ln}_4(\text{Al}_2\text{O}_{7-x}\text{F}_{2x})\text{O}_2$  (top set) ( $\text{Ln} = \text{Sm}, \text{Eu}, \text{Gd}$ ). The relatively low intensity of all the spectra could be a priori attributed to the method of synthesis, where a low preparation temperature was used and thus low crystallinity was expected. The Raman spectra of the starting  $\text{Ln}_4(\text{Al}_2\text{O}_7\Box)\text{O}_2$  materials are quite similar, since they are structurally akin, and are in good agreement with the bibliography (Hasdinator-Bin-Hassan, 2010). An evaluation of the whole spectra is beyond the scope of this work due to the complexity of the structure, so only the high-frequency region will be treated in detail. The as-prepared samples show four well defined bands between 700 and  $800\text{ cm}^{-1}$  that can be unambiguously ascribed to Al—O

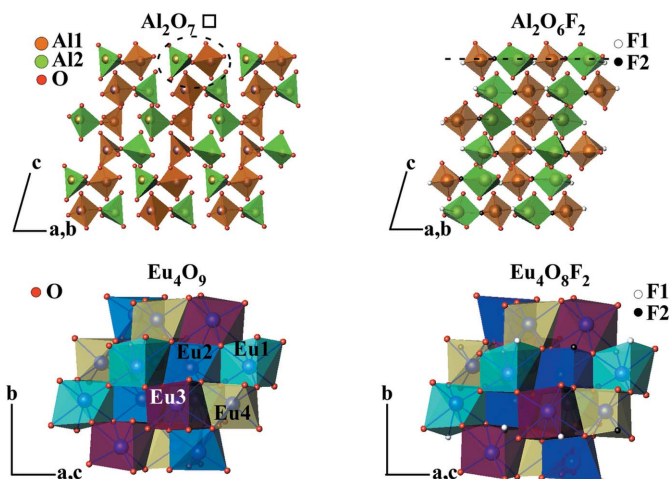


Figure 8

As an example, a polyhedral view of the  $\text{Eu}_4(\text{Al}_2\text{O}_7\Box)\text{O}_2$  and  $\text{Eu}_4(\text{Al}_2\text{O}_6\text{F}_2)\text{O}_2$  phases obtained from the Rietveld refinement structural data using *Atoms62* software.

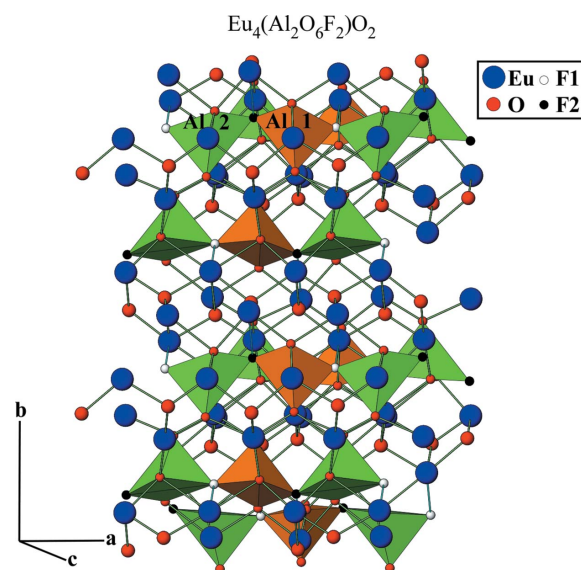


Figure 9

As an example, a simplified representation of the new  $\text{Eu}_4(\text{Al}_2\text{O}_6\text{F}_2)\text{O}_2$  phase structure obtained from the Rietveld refinement structural data using *Atoms62* software.

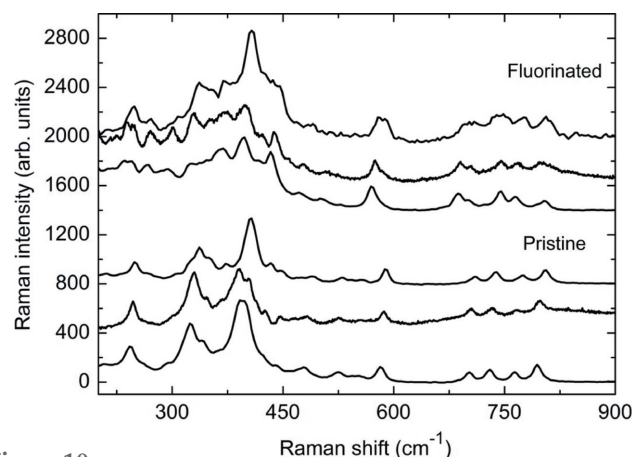


Figure 10

Raman spectra of the pristine  $\text{Ln}_4(\text{Al}_2\text{O}_7\Box)\text{O}_2$  and  $\text{Ln}_4(\text{Al}_2\text{O}_6\text{F}_2)\text{O}_2$  samples. Bottom to top:  $\text{Ln} = \text{Sm}, \text{Eu}, \text{Gd}$ .



stretching modes, since these have shorter bond distances than  $\text{Ln}-\text{O}$ . In the cuspidine structure, the existence of pyroaluminate units of  $[\text{Al}_2\text{O}_7]$  type suggests that it is appropriate to separate the expected modes into internal pyrogroup modes and lattice modes, although the covalent degree of the  $\text{Al}-\text{O}$  bond is lower than that of  $\text{Si}-\text{O}$  or  $\text{P}-\text{O}$  bonds in  $[\text{Si}_2\text{O}_7]$  or  $[\text{P}_2\text{O}_7]$  groups. Moreover, since these units are disconnected within the structure, correlation effects can be dismissed.

Following this approach, and taking into consideration the crystallographic results, the  $[\text{Al}_2\text{O}_7]$  units can be considered as consisting of two  $\text{AlO}_3$  pyramids connected by a bridging oxygen  $\text{O}'$  [O(5) in Tables S2 and S3] in the form  $\text{O}_3-\text{Al}-\text{O}'-\text{Al}-\text{O}_3$ . Within this model, the expected modes can be divided into vibrational modes of the  $\text{AlO}_3$  pyramids and those of the  $\text{Al}-\text{O}'-\text{Al}$  bridge.

A regular pyramid with  $C_{3v}$  symmetry would give two stretching modes in the region of study, one  $A_1$  mode and one  $E$  mode, consisting mainly of the vibration of the three oxygens of the pyramid along the  $\text{Al}-\text{O}$  bonds. However, since Al is located in a  $4e$  site with very low local symmetry ( $C_1$ ), the pyramids must be considered as irregular, giving three  $A$  modes. On the other hand, the  $\text{Al}-\text{O}'-\text{Al}$  bridge is expected to give two stretching modes: one symmetric mode coming mainly from the vibration of Al atoms and one anti-symmetric mode involving Al and O vibrations. The energy of the former will obviously depend on the cation and is found between  $520-560\text{ cm}^{-1}$  in the case of  $[\text{Ge}_2\text{O}_7]$  (Saez-Puche *et al.*, 1992; Hanuza *et al.*, 2011) and  $[\text{Ga}_2\text{O}_7]$  (Kaminskii *et al.*, 2014) and  $620-700\text{ cm}^{-1}$  for  $[\text{Si}_2\text{O}_7]$  (Achary *et al.*, 2017; Lecleach & Gillet, 1990),  $[\text{P}_2\text{O}_7]$  and  $[\text{S}_2\text{O}_7]$  (Kazuo, 2009). In our case, the Al vibration was expected to be around  $600\text{ cm}^{-1}$  and could be tentatively ascribed to the intense band at  $590\text{ cm}^{-1}$ . Therefore, the only mode from the bridge in the high frequency region would be the antisymmetric mode. Although some authors have considered in analogous systems that the  $\text{O}'$  is located in an inversion centre, thus yielding a Raman forbidden or very weak antisymmetric mode (Saez-Puche *et al.*, 1992), the approximation needs the angle of  $X-\text{O}'-X$  to be close to  $180^\circ$  and both  $X-\text{O}'$  distances to be alike. These assumptions seem to be far from our case, where the  $X-\text{O}'-X$  angle is around  $140^\circ$ .

Since only four modes are observed in the high-frequency region of the  $\text{Ln}_4(\text{Al}_2\text{O}_7\Box)\text{O}_2$  sample, the model that best fits our data is that of two irregular but similar pyramids, which would give three stretching modes, connected by an  $\text{Al}-\text{O}'-\text{Al}$  bridge whose antisymmetric mode would supply the required fourth mode.

The model of the isolated  $[\text{Al}_2\text{O}_7]$  units is not valid anymore for the fluorinated samples, where F is proposed to be located in the interstitial positions between these units as well as substituting for the  $\text{O}'$  in the bridge. Thus,  $\text{AlO}_3\text{F}_2$  quasi-square pyramids sharing F vertices form infinite chains along the  $a$  axis (see Fig. 10). By applying the point group  $C_{2v}$  symmetry operations to the constituent atoms of the pyramid (two F and two O atoms in the base and one apical  $\text{O}_{\text{ap}}$ ), five stretching modes are expected in the high frequency region,

considering that all the pyramids are equivalent: three  $A_1$  ( $\text{Al} + \text{O}_{\text{ap}}$ , F, O), one  $B_1$  ( $\text{Al} + \text{F}$ ) and one  $B_2$  ( $\text{Al} + \text{O}$ ). This number of modes is in good agreement with what we observe in the spectra of the fluorinated samples, where five modes are found in the  $650-820\text{ cm}^{-1}$  region. The agreement with the experimental observation suggests that correlation effects, if present, result in almost degenerate modes that remain unresolved because of the spectral broadening. Regarding the symmetrical mode of the  $\text{Al}-\text{O}'-\text{Al}$  bridge in the pristine samples, its position shifts from  $590$  to  $570\text{ cm}^{-1}$  upon fluorination, which would agree with the substitution of the  $\text{O}'$  bridge by F, supporting the assumption from the structural studies that F is located in this site.

Therefore, the Raman measurements are consistent with the crystallographic model proposed for fluorinated  $\text{Ln}_4(\text{Al}_2\text{O}_6\text{F}_2)\text{O}_2$  cuspidines.

#### 4. Conclusions

In summary, new  $\text{Ln}_4(\text{Al}_2\text{O}_6\text{F}_2)\text{O}_2$  ( $\text{Ln} = \text{Sm}, \text{Eu}, \text{Gd}$ ) phases with a cuspidine-related structure have been synthesized using a low-temperature fluorination route, a technique that uses  $\text{Ln}_4(\text{Al}_2\text{O}_7)\text{O}_2$  as the oxide precursor and poly(vinylidene difluoride) as the fluorination agent. The results illustrate the versatility of this fluorination route for the synthesis of new oxide-fluoride systems. The Raman measurements are consistent with the crystallographic model proposed for new fluorinated  $\text{Ln}_4(\text{Al}_2\text{O}_6\text{F}_2)\text{O}_2$  cuspidines: the incorporation of fluorine in the  $\text{Ln}_4(\text{Al}_2\text{O}_7\Box)\text{O}_2$  structure results in Al coordination changes from four to five, which allows the conversion of isolated  $\text{Al}_2\text{O}_7\Box$  groups into infinite chains of distorted square-based pyramids.

#### Funding information

This research was funded by the Ministerio de Economía, Industria y Competitividad (MAT2016-76739-R) (AEI/FEDER, UE), and Departamento de Educación of the Basque Government (IT-630-13). The authors thank SGiker of UPV/EHU for technical and personnel support. A. Morán-Ruiz thanks UPV/EHU for funding.

#### References

- Achary, S. N., Tyagi, A. K., Gupta, M. K., Mittal, R., Bhattacharya, K. & Rao, R. (2017). *J. Alloys Compd.* **702**, 573–584.
- Berry, F. J., Heap, R., Helgason, Ö., Moore, E. A., Shim, S., Slater, P. R. & Thomas, M. F. (2008). *J. Phys. Condens. Matter*, **20**, 215207–215213.
- Clemens, O., Kruk, R., Patterson, E., Loho, C., Reitz, C., Wright, A. J., Knight, K. S., Hahn, H. & Slater, P. R. (2014). *Inorg. Chem.* **53**, 12572–12583.
- Clemens, O. & Slater, P. R. (2013). *Rev. Inorg. Chem.* **33**, 105–117.
- Dae-Min, K., Sang-Ho, L., William, B. A., Kyeong-Beom, K., Yoon-Suk, O. & Sung-Min, L. (2011). *J. Am. Ceram. Soc.* **94**, 3455–3459.
- Ghosh, S. (2015). *Advanced Ceramic Processing*, edited by A. Mohamed, pp. 111–138. London: InTech.
- Hancock, C. A., Herranz, T., Marco, J. F., Berry, F. J. & Slater, P. R. (2012). *J. Solid State Chem.* **186**, 195–203.



- Hanuza, J., Maczka, M., Ptak, M., Lorenc, J., Hermanowicz, K., Becker, P., Bohatý, L. & Kaminskii, A. A. (2011). *J. Raman Spectrosc.* **42**, 782–789.
- Hasdinor-Bin-Hassan, O. (2010). *Oxides with Polyatomic Anions Considered as New Electrolyte Materials for Solid Oxide Fuel Cells*, pp. 48–53. Jülich: Forschungszentrum Jülich GmbH.
- Heap, R., Slater, P. R., Berry, F. J., Helgason, O. & Wright, A. J. (2007). *Solid State Commun.* **141**, 467–470.
- Kaminskii, A. A., Yu, H. H., Wang, J. Y., Zhang, Y. Y., Zhang, H. J., Lux, O., Rhee, H., Eichler, H. J., Hanuza, J., Yoneda, H. & Shirakawa, A. (2014). *Laser Phys.* **24**, 085803–085810.
- Katayama, T., Chikamatsu, A., Kamisaka, H., Kumigashira, H. & Hasegawa, T. (2016). *Appl. Phys. Expr.* **9**, 025801–025804.
- Kawahara, K., Chikamatsu, A., Katayama, T., Onozuka, T., Ogawa, D., Morikawa, K., Ikenaga, E., Hirose, Y., Harayama, I., Sekiba, D., Fukumura, T. & Hasegawa, T. (2017). *CrystEngComm*, **19**, 313–317.
- Kazuo, N. (2009). Editor. *Infrared and Raman Spectra of Inorganic and Coordination Compounds, Part A: Theory and Applications in Inorganic Chemistry*. Hoboken, New Jersey: John Wiley & Sons.
- Kendrick, E., Russ, M. & Slater, P. R. (2008). *Solid State Ionics*, **179**, 819–822.
- Le Cléach, A. & Gillet, P. (1990). *Eur. J. Mineral.* **2**, 43–54.
- Martín-Sedeño, M. C., Losilla, E. R., León-Reina, L., Bruque, S., Marrero-López, D., Núñez, P. & Aranda, M. A. G. (2004). *Chem. Mater.* **16**, 4960–4968.
- Martín-Sedeño, M. C., Marrero-López, D., Losilla, E. R., Bruque, S., Núñez, P. & Aranda, M. A. G. (2006). *J. Solid State Chem.* **179**, 3445–3455.
- Martín-Sedeño, M. C., Marrero-López, D., Losilla, E. R., León-Reina, L., Bruque, S., Núñez, P. & Aranda, M. A. G. (2005). *Chem. Mater.* **17**, 5989–5998.
- Moon, E. J., Choquette, A. K., Huon, A., Kulesa, S. Z., Barbash, D. & May, S. J. (2015). *APL Mater.* **3**, 062511–062517.
- Morán-Ruiz, A., Vidal, K., Larrañaga, A. & Arriortua, M. I. (2018). *Ceram. Int.* **44**, 8761–8767.
- Rodríguez-Carvajal, J. (2010). *FULLPROF*. Version 4.80. ILL, Grenoble, France.
- Saez-Puche, R., Bijkerk, M., Fernández, F., Baran, E. J. & Botto, I. L. (1992). *J. Alloys Compd.* **184**, 25–34.
- Shape Software (2005). *ATOMS*, version 6.2. Shape Software, Kingsport, Tennessee, USA.
- Slater, P. R. (2002). *J. Fluor. Chem.* **117**, 43–45.
- Zhou, X., Xu, Z., Fan, X., Zhao, S., Cao, X. & He, L. (2014). *Mater. Lett.* **134**, 146–148.